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(11) **EP 0 765 951 B1**

(12)

EUROPEAN PATENT SPECIFICATION

(45) Date of publication and mention
of the grant of the patent:
06.12.2000 Bulletin 2000/49

(51) Int Cl.7: **C23C 4/02, F01D 11/12**

(21) Application number: **96307035.4**

(22) Date of filing: **26.09.1996**

(54) **Abradable ceramic coating**

Verschleissfeste keramische Beschichtung

Revêtement céramique résistant

(84) Designated Contracting States:
DE ES FR GB

(30) Priority: **26.09.1995 US 534146**

(43) Date of publication of application:
02.04.1997 Bulletin 1997/14

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Description

[0001] This invention relates generally to duct segments for use in gas turbine engines, and particularly to ceramic abradable coatings for such duct segments.

[0002] Modern gas turbine engines, particularly those used in aircraft, operate at high rotational speeds and high temperatures for increased performance and efficiency. The turbine of a modern gas turbine engine is typically of an axial flow design and includes a plurality of axial flow stages. Each axial flow stage comprises a plurality of blades mounted radially at the periphery of a disk which is secured to a shaft. A plurality of duct segments surround the stages to limit the leakage of gas flow around the tips of the blades. These duct segments are located on the inner surface of a static housing or casing. The incorporation of the duct segments improves thermal efficiency because more work may be extracted from gas flowing through the stages as opposed to leaking around the blade tips.

[0003] Although the duct segments limit the leakage of gas flow around the blade tips, they do not completely eliminate the leakage. It has been found that even minor amounts of gas flow around the blade tips detrimentally affect turbine efficiency. Thus, gas turbine engine designers proceed to great lengths to devise effective sealing structures. These structures generally include a coated duct segment in combination with a blade tip coating which renders the tips resistant to wear. In operation, the tips provide sealing by cutting into the coating on the duct segment.

[0004] Unfortunately current duct segment coatings, which are typically ceramic, suffer from excessive material loss as a result of erosion or spalling. In general, erosion is the wearing away of coating material due to factors such as abrasion and corrosion. Erosion often results from particle impingement during engine operation. Spalling is typically caused by delamination cracking at the ceramic-metal interface resulting from thermal stress and the aggressive thermal environment. Spalling is essentially piecemeal coating loss consisting of many small coherent volumes of coating material. Ceramic coating loss increases blade tip clearance and thus is detrimental to turbine efficiency, as well as detrimental to the blades themselves. For example, the blades may become damaged due to the increased temperature at which the engine must then operate to make up for lost thrust.

[0005] Accordingly, there exists a need for a coating which is abradable as well as erosion and spalling resistant. This coating is necessary for a sealing system having superior abradability and erosion resistance.

[0006] The present invention is directed towards providing a coating which is abradable as well as erosion and spalling resistant.

[0007] From one aspect, the invention provides a segmented abradable ceramic coating system comprising a segmented abradable ceramic layer and a MCrAlY bondcoated substrate, said abradable ceramic layer comprising a material selected from the group consisting of a material having a compositional blend of zirconia partially stabilized with yttria and zirconia fully stabilized with yttria; a material having a compositional blend of zirconia partially stabilized with ceria and zirconia fully stabilized with ceria; a material having a compositional blend of zirconia partially stabilized with magnesia and zirconia fully stabilized with magnesia; and a material having a compositional blend of zirconia partially stabilized with calcia and zirconia fully stabilized with calcia, wherein the abradable ceramic layer is located on said MCrAlY bondcoated metallic substrate, the layer consisting essentially of powder particles having a spherical and hollow morphology prior to deposition for increased abradability.

[0008] A further aspect of the invention includes a segmented abradable ceramic coating system having enhanced abradability. The system comprises a duct segment including a metallic substrate; a MCrAlY bond coat on the substrate; and a segmented abradable ceramic (SAC) coating on the MCrAlY bond coat. The nature of the MCrAlY bond coat is such that it must provide sufficient resistance to oxidation and corrosion. One aspect of the SAC coating comprises three ceramic layers which include a base coat foundation layer of material selected from the group consisting of zirconia stabilized with ceria, zirconia stabilized with magnesia, zirconia stabilized with calcia, zirconia stabilized with yttria, and mixtures thereof; an abradable top layer comprising zirconia; and a graded interlayer which is a compositional blend of the base coat foundation layer and the abradable top layer. The graded interlayer is positioned between the base coat foundation layer and the abradable top layer. The segmented abradable ceramic coating also includes a plurality of vertical microcracks and the three ceramic layers comprise powder particles which are spherical and hollow prior to deposition for increased abradability.

[0009] The segmented abradable sealing system has enhanced abradability relative to segmented sealing systems known in the art. In one preferred embodiment, the sealing system comprises a duct segment including a metallic substrate; a MCrAlY bond coat on the substrate; and a segmented abradable ceramic coating on the MCrAlY bond coat. The system also includes a cooperating interacting turbine component having an abrasive coating on a portion thereon such that the abrasive coating can interact with the segmented abradable ceramic coating to provide sealing.

[0010] Yet another broad aspect of the invention includes a method of making a segmented abradable ceramic coating. By precisely controlling the deposition parameters and utilizing specific powder compositions and powder morphology, segmentation of the coating into vertical microcracks is achieved as well as superior abradability and erosion resistance.

[0011] An advantage of the present invention in its preferred embodiments at least is segmentation of the coating

into columnar type cells thereby significantly improving ceramic spalling resistance. The plasma spray processing parameters disclosed herein produce ceramic segmentation which enhances erosion resistance and results in superior abrasability.

[0012] Fig. 1 shows a cross-section of a portion of a modern gas turbine engine.

[0013] Fig. 2 is a flow path duct segment incorporating the present invention.

[0014] As shown in Fig. 1, a plurality of blades 2 mounted on a disk rotate about an axis of rotation 4 in the turbine section of a gas turbine engine. A static housing, concentric with the axis of rotation 4, surrounds the blades 2. A gap 8 exists between the housing and tips 11 of blades 2.

[0015] During gas turbine engine operation, high temperature gas flows between the rotating turbine blades 2. The turbine must efficiently convert the energy of this high temperature gas into shaft horsepower to drive a compressor. Thus, gas leakage through gap 8 must be minimized because minute changes in tip clearance have a great adverse effect on gas turbine engine performance.

[0016] Accordingly, a flow path duct segment 10 is provided between the blade tips 11 and the housing. As shown in Fig. 2, the flow path duct segment 10 includes a substrate or seal shoe 12 which is made of a nickel or cobalt base superalloy and is typically fabricated by casting and machining. The substrate 12 is located on the inner wall of the housing.

[0017] An abradable ceramic coating system is applied to the substrate 12. Prior to depositing the coating system, the substrate 12 may be cleaned to remove contamination. Cleaning is conventional and may include aluminum oxide grit blasting.

[0018] A bond coat 14 of an MCrAlY material is applied to the substrate 12. MCrAlY refers to known metal coating systems in which M denotes nickel, cobalt, iron, or mixtures thereof; Cr denotes chromium; Al denotes aluminum; and Y denotes yttrium. MCrAlY materials are often known as overlay coatings because they are put down in a predetermined composition and do not interact significantly with the substrate during the deposition process. For examples of MCrAlY materials see U.S. Patent No. 3,528,861 which describes a FeCrAlY coating as does U.S. Patent No. 3,542,530. In addition, U.S. Patent No. 3,649,225 describes a composite coating in which a layer of chromium is applied to a substrate prior to the deposition of a MCrAlY coating. U. S. Patent No. 3,676,085 describes a CoCrAlY overlay coating while U.S. Patent No 3,754,903 describes a NiCoCrAlY overlay coating having particularly high ductility. U.S. Patent No. 4,078,922 describes a cobalt base structural alloy which derives improved oxidation resistance by virtue of the presence of a combination of hafnium and yttrium. A preferred MCrAlY bond coat composition is described in U.S. Patent No. Re. 32,121 as having a weight percent compositional range of 5-40 Cr, 8-35 Al, 0.1-2.0 Y, 0.1-7 Si, 0.1-2.0 Hf, balance selected from the group consisting of Ni, Co and mixtures thereof. See also U.S. Patent No. 4,585,481.

[0019] This MCrAlY bond coat 14 may be applied by any method capable of producing a dense, uniform, adherent coating of desired composition. For example, techniques such as sputtering, electron beam physical vapor deposition and high velocity plasma spray techniques are known. In the latter technique, a spray torch may operate in a vacuum chamber at a pressure of less than about 60 torr (60 mm Hg) or in another suitable atmosphere, such as air. If a vacuum chamber is employed, the substrate is heated to a temperature between about 1500°F (816°C) and about 1900°F (1038°C). If an air atmosphere is used, the substrate temperature is maintained at less than about 600°F (316°C). Preferably, however, the bond coat is applied by a process known as high velocity oxy-fuel (HVOF) spray. This deposition process utilizes a spray torch in which liquid fuel or gas is combusted with oxygen to produce a high velocity gas stream into which powdered coating material is injected, heated and propelled onto the part. This process is effective as well as cost efficient.

[0020] The particle size for the bond coat 14 may be between about 15 microns (.015 mm) and about 60 microns (.060 mm), with preferably a mean particle size of about 25 microns (.025 mm). The bond coat may be applied to a thickness between about 5 mils (.127 mm) and about 10 mils (.254 mm). Preferably the thickness is between about 6 mils (.152 mm) and about 7 mils (.178 mm).

[0021] Next a segmented abradable ceramic (SAC) coating 16 is applied on bond coat 14. The SAC coating 16 comprises three ceramic layers which are individually applied for an overall thickness between about 20 mils (.508 mm) and about 75 mils (1.905 mm), preferably about 50 mils (1.270 mm). The SAC coating 16 is typically produced in one continuous spray process. However, three separate spray events may be employed.

[0022] In a suitable setup for deposition of the above described layers, a plurality of bondcoated substrates 12 are loaded into a hollow cylindrical fixture such that the bondcoated surfaces face the inner diameter of the cylindrical fixture. A plasma spray gun is positioned in the interior of the cylindrical fixture for depositing the layers.

[0023] First, a base coat foundation layer 18 is applied to bond coat 14 to a thickness of between about 5 mils (.127 mm) and about 15 mils (.381 mm). Layer 18 is preferably a yttria partially stabilized zirconia ceramic layer (yttria partially stabilized zirconia herein refers to a composition of about 12 weight percent or less yttria stabilized zirconia). However, a composition of between about 6 weight percent and about 20 weight percent yttria stabilized zirconia may be used, with a preferred range between about 7 weight percent and about 12 weight percent yttria stabilized zirconia for material

strength. Similarly, other zirconia based compositions, such as ceria stabilized zirconia, magnesia stabilized zirconia, calcia stabilized zirconia and mixtures thereof may be substituted for the yttria stabilized zirconia. A blended layer having a combination of about 7 weight percent yttria stabilized zirconia and other yttria stabilized zirconia powders may also be employed.

5 **[0024]** The particle size of the powder used for layer 18 (as well as the powder used for subsequently applied layer 20 and layer 22) may range from about 5 microns (.005 mm) to about 175 microns (.175 mm) with a preferred particle size of about 50 microns (.050 mm) in mean diameter. The particles for layer 18, as well as for layers 20 and 22, are produced from a spray dried and sintered process which results in spherical and hollow powders, as opposed to fused and crushed powders which are angular and solid. In general, the initial step of the spray dried and sintered process
10 includes mixing raw zirconia and yttria to a desired weight percent ratio. This mixture is then combined with water (and conventional binders) to produce a slip. The slip is then fed into a spray dryer which partially dries the slip by spraying the material into a heated chamber, thereby producing spherical and hollow powders. Then the material is heated at the sintering temperature in a furnace for typically between about 4 hours and about 8 hours. This sintering temperature therein is usually about 60% to about 70% of the theoretical melting point of zirconium oxide.

15 **[0025]** Alternatively, a spray dried and plasma densified process may be used, although this process may be more expensive than the above described process. In general, the initial step of this process also includes mixing raw zirconia and yttria to a desired weight percent ratio. This mixture is then also combined with water (and conventional binders) to produce a slip. The slip is then fed into a spray dryer which partially dries the slip by spraying the material into a heated chamber, thereby producing spherical and hollow powder. Following the spray drying step, however, the powder
20 is then fed through a plasma spray gun where the yttria and zirconia melt to produce a homogeneous composition.

[0026] The spherical and hollow morphology of the powder prior to deposition is a key factor for the success of the present invention, especially with respect to its superior abrasability. For example, if solid particles are present in the powder, more heat is required to melt the powder. This results in a dense coating which may not be very abrasable. Also, the deposition efficiency for angular and solid particles is less relative to spherical and hollow particles. This is
25 extremely important for manufacturing cost concerns.

[0027] Layer 18, the base coat foundation layer, is beneficial to the success of the segmented abrasable coating system because it provides a tough ceramic structure, starts segmentation of the deposited material into vertical microcracks, provides erosion protection and provides a thermal barrier benefit. In addition, layer 18 bonds to the MCrAlY bond coat 14.

30 **[0028]** Layer 18 is typically plasma sprayed in air. It may be desirable to heat the substrate 12 and monitor the temperature at less than about 600°F (316°C) to help segmentation of the material into vertical microcracks. This heating may be accomplished by application of heat to the back side of the substrate 12 during material deposition. The above described heating parameters apply to layer 18, as well as to the other subsequently applied layers. Preferably, however, the substrate 12 is not heated except incidentally during spraying.

35 **[0029]** The processing parameters of the present invention are controlled to produce vertical segmentation (approximately perpendicular to the bond coat surface) and are specific to variables such as gun type and fixture geometry. In general, we have found that a close gun to part spray distance coupled with relatively high power deposition results in desirable vertical segmentation of between about 4 and about 8 microcracks per inch. The parameters described herein were specifically tailored for use with a Sulzer Metco, Inc. 3MB air plasma spray gun and a cylindrical fixture
40 having a 30 inch (.76m) diameter. One of ordinary skill in the art would appreciate that the parameters may vary with the use of a different spray gun and/or fixture. Accordingly, the parameters set forth herein may be used as a guide for selecting other suitable parameters for different operating conditions.

[0030] Specifically, during the spray deposition of layer 18, the cylindrical fixture rotates at a speed between about 5 rpm and about 25 rpm, and preferably at about 12 rpm. The plasma spray gun is located in the interior of the hollow
45 cylindrical fixture. The gun to part angle during individual part coating is between about 80 degrees and about 100 degrees, and preferably about 90 degrees. The gun to part distance is varied in increments from about a nominal 2 inches (.05m) (starting distance) to about a nominal 5 inches (.13m) (end distance), and preferably between about 2.75 inches (.07m) (starting distance) and about 3.25 inches (.083m) (end distance) during production of layer 18. This close gun distance is necessary for satisfactory vertical segmentation. Gun traverse speed across each part during
50 deposition is between about 1 in/min (.03m/min) and about 5 in/min (.13m/min), preferably about 4.4 in/min (.11m/min).

[0031] Powder feed rate is between about 15 grams/min and about 50 grams/min, and preferably about 35 grams/min. Carrier gas flow, such as nitrogen, is used to maintain the powder under pressure and facilitate powder feed. The flow rate is between about 5 scfh (standard cubic feet/hour) (.14 scmh (standard cubic meters/hour) and about 20 scfh (.57 scmh), preferably about 11 scfh (.31 scmh). Standard conditions are herein defined as about room temperature
55 (25°C) and about one atmosphere of pressure (101 kPa). Primary gas flow, such as nitrogen gas, in the gun is between about 80 scfh (2.27 scmh) and about 120 scfh (3.40 scmh), and preferably about 99 scfh (2.80 scmh). Similarly, secondary gas flow, such as hydrogen, in the gun is between about 5 scfh (.14 scmh) and about 30 scfh (.85 scmh), and preferably about 18 scfh (.51 scmh). Gun voltage is between about 60 volts and about 80 volts, and preferably about

75 volts. Similarly, gun amperage is between about 700 amps and about 900 amps, and preferably about 736 amps. We have found the above described parameters to be optimum for the deposition process using the Sulzer Metco 3MB plasma spray gun, but one skilled in the art would appreciate that the parameters are dependent on variables, including but not limited to, powder type, powder size and especially type of gun.

[0032] Next a graded interlayer 20 is applied to base coat foundation layer 18 to a thickness between about 3 mils (.076 mm) and about 10 mils (.254 mm). This layer is also typically plasma sprayed in air. The composition of the graded interlayer 20 is a blend of layer 18 (base coat foundation layer) and layer 22, which is an abradable top coat subsequently applied to layer 20. For ease of describing the composition of graded interlayer 20, the composition of layer 22 will now be described. Layer 22 is an abradable top layer comprising zirconia. The nature of layer 22 is such that it must be soft enough to allow blade tips to cut into layer 22 and provide sealing. The composition of layer 22 is typically a blend of 7 weight percent yttria stabilized zirconia and 20 weight percent yttria stabilized zirconia. The ratio of the blend depends on the desired characteristics of the resulting deposit. For example, if increased erosion resistance is desired, then an increase in the amount of 7 weight percent yttria stabilized zirconia should be employed, whereas if an increase in abradability is desired, then more 20 weight percent yttria stabilized zirconia should be added. However, in another embodiment essentially 100% of zirconia fully stabilized with yttria, such as 20 weight percent yttria stabilized zirconia, may be employed for layer 22.

[0033] The composition of graded interlayer 20 is typically varied from a starting composition of a weight percent ratio of layer 18/layer 22 to a final composition of a weight percent ratio of layer 18/layer 22. For example, we have employed a starting composition of 90/10 (weight percent ratio of layer 18 to layer 22) to a final composition of 10/90 (weight percent ratio of layer 18 to layer 22).

[0034] The deposition parameters for production of layer 20 are the same as those previously described for production of layer 18, except that in this case, preferably the gun to part distance is held constant at about 3.25 inches (.083 m). It may be possible to vary the gun to part distance within the ranges described for production of layer 18, but we have found that keeping the distance constant is optimum. The advantage of this graded interlayer is that it provides a strength link between layer 18 and layer 22.

[0035] It should also be noted that the graded layer 20 may also be produced by other means, such as by application of individual layers of layer 18 and layer 22, varied by pass.

[0036] After application of the graded layer 20, the above described abradable layer 22 is sprayed on the graded interlayer 20 to a thickness between about 15 mils (.381 mm) and about 55 mils (1.397 mm). Preferably, the thickness of layer 22 is about 35 mils (.889 mm). The deposition parameters for production of layer 22 are the same as those described for production of layer 20.

[0037] Porosity may be intentionally created within layer 22 by adding small amounts of materials such as polyester or Lucite® powder. The inclusion of 1 to about 7 weight percent polyester powder (60 micron (.060 mm) nominal particle size) in layer 22 may produce a porosity on the order of about 20-30 volume percent. High porosity levels, such as levels greater than about 25 volume percent, may be unsatisfactory because of potential erosion of the coating. However, since a dense structure is desired for erosion resistance, addition of these materials should be minimized, if not eliminated entirely. It is desirable for the resultant density of layer 22 to be between about 90-95 percent theoretical.

[0038] In an alternative embodiment of the present invention, layer 22 may comprise alternating layers of a layer of 20 weight percent yttria stabilized zirconia and a layer of blended 7 weight percent yttria stabilized zirconia and 20 weight percent yttria stabilized zirconia (such as a 50-50 blend) for an overall thickness of preferably about 35 mils (.889 mm). The thickness of the layers may be between about .5 mils (.013 mm) and about 5 mils (.127 mm) each. Each layer should be about the same thickness.

[0039] In another embodiment of the invention, other materials, including but not limited to ceria, magnesia, calcia or mixtures thereof may be employed in place of yttria for the SAC system. However, yttria stabilized zirconia materials are recommended for SAC applications exceeding 1950°F (1066°C).

[0040] In another embodiment of the invention, alumina (99.0% purity) may be employed. For example, a thin layer (less than about 5 mils (.127 mm)) of alumina may be sprayed on the bond coat 14 prior to application of layer 18. Alternatively, a blended composition of alumina and less than about 12 weight percent yttria stabilized zirconia may be used for the base coat foundation layer 18 (less than 10 weight percent alumina blend). The thin alumina layer may also be applied upon completion of layer 18 and prior to application of layer 20.

[0041] In yet another embodiment of the invention, the SAC coating may consist essentially of layer 22. This coating would be desirable for military applications. Due to the smaller size of military gas turbine engine components, a thinner SAC coating is acceptable.

[0042] After application of the SAC coating, the flow path duct segment 10 is usually heat treated for stress relief. Specifically, the duct segment 10 may be heat treated at about 1975°F (1079°C) +/- 25°F (14°C), for about 4 hours and then force cooled to about 1100°F (593°C) in about 22 minutes or less. The segment 10 is then force cooled to about 1000°F (538°C) in about 7 minutes or less and again force cooled to below 300°F (149°C). This heat treatment preserves or even increases the useful life of the duct segment 10. For example, spallation of the coating is reduced.

This heat treatment may also be employed after deposition of layer 14 (bond coat) and prior to deposition of the foundation layer 18, although it is not necessary for practice of the present invention.

[0043] The above described coating system is particularly suited to interact with blade tips which are coated on their free end with an abrasive material, including but not limited to cubic boron nitride. This interaction provides an effective sealing system.

[0044] The present invention will now be described by example which is meant to be exemplary rather than limiting.

Example

[0045] Nickel based high pressure turbine duct segments which were previously coated with a NiCoCrAlY oxidation resistant bondcoat were loaded into a hollow cylindrical fixture having a 30 inch (.76m) diameter such that the bond coated surface of the duct segments faced the center of the fixture. The HVOF process previously described was used to apply the NiCoCrAlY bondcoat to the duct segments.

[0046] A Sulzer Metco, Inc. 3MB air plasma spray gun was positioned in the interior of the fixture which rotated at about 12 rpm. The gun was located at about a 90 degree angle to the inside surface of each duct segment to be coated in turn. Three distinct layers were created on the bondcoated duct segments using the parameters set forth in Table 1 below. The layers were deposited in one continuous spray process to avoid any potential weak links in the overall coating due to momentarily stopping the process. This continuous process was possible with the use of several Miller Thermal Model 1250 powder feeders, each containing powder having a specific composition. The powders also had a spherical and hollow morphology prior to deposition. The feeders can be computer controlled to deposit the desired composition for each layer.

[0047] Specifically, layer 1, a base coat foundation layer, 0.010 inches (.254 mm) to 0.015 inches (.381 mm) thick of between about 6 weight percent and about 8 weight percent yttrium oxide, remainder zirconium oxide (Sulzer Metco, Inc. 204NS powder) was deposited. Next, layer 2, a graded interlayer of varied composition from 90 weight percent layer 1 / 10 weight percent layer 3 (starting composition) to 10 weight percent layer 1 / 90 weight percent layer 3 (end composition) was deposited. The grading was performed in 8 uniform steps over a layer thickness of 0.005 inches (.013 mm) at a constant gun distance of 3.25 inches (.083m), as noted in Table 1 below.

[0048] Layer 3, an abradable top layer, was then deposited to a thickness between about 0.025 inches (.635 mm) and 0.040 inches (1.016 mm). The composition of layer 3 was a 50-50 blend of 1) the composition of layer 1 and 2) between about 18.5 weight percent and about 21.50 weight percent yttrium oxide, remainder zirconium oxide (Sulzer Metco, Inc. 202 powder or equivalent). A multi-layer segmented abradable coating having between about 4 and about 8 vertical microcracks per inch resulted.

Table 1

	Layer 1	Layer 2	Layer 3
fixture rotation speed (rpm)	12	12	12
gun traverse speed across ea. seal (in/min) [m/hr]	4.4 [6.7]	4.4 [6.7]	4.4 [6.7]
gun to seal distance (in) [m]			
start	2.75 [.07]	3.25 [.083]	3.25 [.083]
end	3.25 [.083]	3.25 [.083]	3.25 [.083]
gun to seal angle (degrees)	90	90	90
gun voltage (volts)	75	75	75
primary gas flow, N ₂ (scfh) [scmh]	99 [2.80]	99 [2.80]	99 [2.80]
secondary gas flow, H ₂ (scfh) [scmh]	18 [.51]	18 [.51]	18 [.51]
gun amperage (amps)	736	736	736
carrier gas flow, N ₂ (scfh) [scmh]	11 [.31]	11 [.31]	11 [.31]
powder feed rate (grams/min)	35	35	35

[0049] An advantage of the present invention is its superior abradability as compared to a segmented coating having a composition of 7 weight percent yttria, balance substantially zirconia (7 YSZ). This superior abradability of the present invention results, in part, because 7 YSZ contains approximately equivalent proportions of tetragonal ZrO₂ and cubic ZrO₂. Tetragonal ZrO₂ is stronger than cubic ZrO₂ due to its lattice structure. Thus, 7 YSZ is a strong, effective erosion

resistant material. However, 20 weight percent yttria, balance substantially zirconia (20 YSZ) does not contain tetragonal ZrO_2 (only cubic ZrO_2), and thus is more abrasible (i.e. softer). As stated previously, 20 YSZ is a desirable composition for the abrasible top layer 22. In a duct segment application, effective abrasibility is an essential requirement for efficient engine operation. Different material combinations can be selected for specific engine models having different needs. For example, a SAC coating with less abrasibility and more erosion resistance can be selected, if desired. In such a case, SAC coating may comprise a blend of 7 YSZ and 20 YSZ as the abrasible layer (layer 22). The blend would provide the needed erosion resistance (from the 7 YSZ) without sacrificing abrasibility. An important point to remember is that the segmented abrasible coating of the present invention, with its unique structure and variety of material combinations, can be tailored for engine specific duct segment applications. The three layer approach provides a means of tailoring the long-term thermal insulation benefit provided by the initial layers (layer 18 and layer 20) and the abrasibility benefit provided by the top layer (layer 22) in a given engine application.

[0050] Although the invention has been shown and described with respect to detailed embodiments thereof, it should be understood by those skilled in the art that various changes in form and detail may be made without departing from the [0051] scope of the invention. Specifically, although the present invention has been described as a segmented abrasible ceramic coating for aircraft gas turbine engine duct segments, the present invention may have other potential applications, including but not limited to, a thermal barrier coating on gas turbine engine components, such as vanes, and as a segmented abrasible ceramic coating system for land based turbine applications. The present invention may also have application in the automotive industry as a coating for automotive engine components, such as pistons.

Claims

1. A segmented abrasible ceramic coating system comprising:

a metallic substrate (12);
a MCrAlY bond coat (14) on the substrate; and
a segmented abrasible ceramic coating (16) on the MCrAlY bond coat, said segmented abrasible ceramic coating comprising three ceramic layers which include a base coat foundation layer (18) comprising a layer of material selected from the group consisting of zirconia stabilized with ceria, zirconia stabilized with magnesia, zirconia stabilized with calcia, zirconia stabilized with yttria, and mixtures thereof; an abrasible top layer (22) comprising zirconia; and a graded interlayer (20) which is a compositional blend of the base coat foundation layer and the abrasible top layer, said graded interlayer positioned between the base coat foundation layer and the abrasible top layer, said segmented abrasible ceramic coating having a plurality of vertical microcracks, wherein the three ceramic layers consist essentially of powder particles which are spherical and hollow prior to deposition for increased abrasibility.

2. The system of claim 1 wherein the base coat foundation layer (18) is a ceramic layer between about 5 mils (.127 mm) and about 15 mils (.381 mm) thick, said base coat foundation layer selected from the group consisting of a material having a composition less than about 12 weight percent ceria stabilized zirconia, a material having a composition less than about 12 weight percent magnesia stabilized zirconia, a material having a composition less than about 12 weight percent calcia stabilized zirconia, a material having a composition less than about 12 weight percent yttria stabilized zirconia, and mixtures thereof.

3. The system of claim 1 wherein the base coat foundation layer (18) comprises a compositional blend of alumina and less than about 12 weight percent yttria stabilized zirconia.

4. The system of claim 1, 2 or 3 wherein the abrasible top layer (22) is a ceramic layer between about 15 mils (.381 mm) and about 55 mils (1.397 mm) thick, said abrasible top layer comprising a blend of zirconia partially stabilized with ceria and zirconia fully stabilized with ceria.

5. The system of 1, 2 or 3 wherein the abrasible top layer (22) is a ceramic layer between about 15 mils (.381 mm) and about 55 mils (1.397 mm) thick, said abrasible top layer comprising a blend of zirconia partially stabilized with magnesia and zirconia fully stabilized with magnesia.

6. The system of 1, 2 or 3 wherein the abrasible top layer (22) is a ceramic layer between about 15 mils (.381 mm) and about 55 mils (1.397 mm) thick, said abrasible top layer comprising a blend of zirconia partially stabilized with yttria and zirconia fully stabilized with yttria.

7. The system of claim 1,2 or 3 wherein the abrasible top layer (22) comprises a material selected from the group consisting of zirconia fully stabilized with ceria, zirconia fully stabilized with magnesia, zirconia fully stabilized with calcia, and zirconia fully stabilized with yttria.
- 5 8. The system of claim 7 wherein the abrasible top layer (22) has a composition of substantially 100 weight percent zirconia fully stabilized with yttria.
9. The system of claim 1, 2 or 3 wherein the abrasible top layer (22) comprises a blend of 7 weight percent yttria stabilized zirconia and 20 weight percent yttria stabilized zirconia.
- 10 10. The system of claim 1 wherein
the base coat foundation layer is between about 5 mils (.127 mm) and about 15 mils (.381 mm) thick, said
base coat foundation layer comprising yttria stabilized zirconia, between about 7 weight percent and about 12
15 weight percent, with a particle size of about 50 microns (.050 mm) in mean diameter;
the graded interlayer is between about 3 mils (.076 mm) and about 10 mils (.254 mm) thick, said graded
interlayer comprising a compositional blend of the base coat foundation layer and the abrasible top layer; and
the abrasible top layer is between about 15 mils (.381 mm) and about 55 mils (1.397 mm) thick, said abrasible
top layer comprising 20 weight percent yttria stabilized zirconia.
- 20 11. A duct segment comprising the segmented abrasible ceramic coating system as claimed in any preceding claim.
12. A sealing system comprising a duct segment as claimed in claim 11 and a cooperating interacting turbine compo-
nent having an abrasive coating on a portion thereon such that said abrasive coating can interact with said seg-
25 mented abrasible coating to provide sealing.
13. A method of producing a segmented abrasible ceramic coating including a base coat foundation layer (18), a
graded interlayer (20), and an abrasible top layer (22), said method comprising:
30 applying the base coat foundation layer (18) on a MCrAlY bondcoated metallic substrate (12) using a spray
gun which heats and propels powdered coating material onto said metallic substrate (12), said base coat
foundation layer (18) comprising a layer of material selected from the group consisting of zirconia stabilized
with ceria, zirconia stabilized with magnesia, zirconia stabilized with calcia, zirconia stabilized with yttria, and
35 mixtures thereof, wherein the distance between the gun and surface to be coated is varied during production
of the base coat foundation layer,
applying the graded interlayer (20) on the base coat foundation layer (18) using a spray gun which heats and
propels powdered coating material onto said base coat foundation layer (18), said graded interlayer comprising
a compositional blend of the base coat foundation layer and the abrasible top layer; wherein the distance
40 between the gun and surface to be coated is held constant during production of the graded layer, and
applying the abrasible top layer (22) on the graded interlayer (20) using a spray gun which heats and propels
powdered coating material onto said graded interlayer (20), said abrasible top layer comprising zirconia,
wherein the distance between the gun and surface to be coated is held constant during production of the
abrasible top layer, wherein each layer consists essentially of powder particles which are spherical and hollow
45 prior to deposition for increased abrasibility.
14. The method of claim 13 wherein the base coat foundation layer is produced to a thickness between about 5 mils
(.127 mm) and about 15 mils (.381 mm) and the graded layer to a thickness between about 3 mils (.076 mm) and
about 10 mils (.254 mm).
- 50 15. A segmented abrasible ceramic coating system consisting essentially of a segmented abrasible ceramic layer
and a MCrAlY bondcoated substrate, said abrasible ceramic layer comprising a material selected from the group
consisting of a material having a compositional blend of zirconia partially stabilized with yttria and zirconia fully
stabilized with yttria; a material having a compositional blend of zirconia partially stabilized with ceria and zirconia
fully stabilized with ceria; a material having a compositional blend of zirconia partially stabilized with magnesia and
55 zirconia fully stabilized with magnesia; and a material having a compositional blend of zirconia partially stabilized
with calcia and zirconia fully stabilized with calcia, wherein the abrasible ceramic layer is located on said MCrAlY
bondcoated metallic substrate, the layer consisting essentially of powder particles having a spherical and hollow
morphology prior to deposition for increased abrasibility.

16. The coating system of claim 15 wherein the ceramic layer is between about 15 mils (.381 mm) and about 55 mils (1.397 mm) thick.

5 Patentansprüche

1. Segmentiertes abreibbares keramisches Beschichtungssystem aufweisend:

ein metallisches Substrat (12),
 10 eine MCrAlY-Bindungsbeschichtung (14) auf dem Substrat, und eine segmentierte abreibbare keramische Beschichtung (16) auf der MCrAlY-Bindungsbeschichtung,
 wobei die segmentierte abreibbare keramische Beschichtung drei keramische Schichten aufweist, zu denen eine Basisbeschichtung-Grundlagenschicht (18), die eine Schicht aus einem Material aufweist, das ausge-
 wählt ist aus der Gruppe, die besteht aus mit Cerdioxid stabilisiertem Zirkoniumdioxid, mit Magnesiumoxid
 15 stabilisiertem Zirkoniumdioxid, mit Calciumoxid stabilisiertem Zirkoniumdioxid, mit Yttriumoxid stabilisiertem Zirkoniumdioxid, und Gemischen davon, eine abreibbare Deckschicht (22), die Zirkoniumdioxid aufweist, und eine Varianz-Zwischenschicht (20), die zusammensetzungsmäßig eine Mischung aus der Basisbeschichtung-Grundlagenschicht und der abreibbaren Deckschicht ist, gehören, wobei die Varianz-Zwischenschicht zwischen der Basisbeschichtung-Grundlagenschicht und der abreibbaren Deckschicht angeordnet ist, wobei die
 20 segmentierte abreibbare keramische Beschichtung eine Mehrzahl vertikaler Mikrorisse hat, und wobei die drei keramischen Schichten im wesentlichen aus Pulverteilchen bestehen, die zwecks erhöhter Abreibbarkeit vor der Abscheidung kugelförmig und hohl sind.

2. System nach Anspruch 1, bei dem die Basisbeschichtung-Grundlagenschicht (18) eine zwischen etwa 5 mil (0,127
 25 mm) und etwa 15 mil (0,381 mm) dicke keramische Schicht ist, wobei die Basisbeschichtung-Grundlagenschicht ausgewählt ist aus der Gruppe, die besteht aus einem Material mit einer Zusammensetzung aus mit weniger als etwa 12 Gew.-% Cerdioxid stabilisiertem Zirkoniumdioxid, einem Material mit einer Zusammensetzung aus mit weniger als etwa 12 Gew.-% Magnesiumoxid stabilisiertem Zirkoniumdioxid, einem Material mit einer Zusammen-
 setzung aus mit weniger als etwa 12 Gew.-% Calciumoxid stabilisiertem Zirkoniumdioxid, einem Material mit einer
 30 Zusammensetzung aus mit weniger als etwa 12 Gew.-% Yttriumoxid stabilisiertem Zirkoniumdioxid, und Gemischen davon.

3. System nach Anspruch 1, bei dem die Basisbeschichtung-Grundlagenschicht (18) ein Zusammensetzungsgemisch aus Aluminiumoxid und mit weniger als etwa 12 Gew.-% Yttriumoxid stabilisiertem Zirkoniumdioxid aufweist.
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4. System nach Anspruch 1, 2 oder 3, bei dem die abreibbare Deckschicht (22) eine zwischen etwa 15 mil (0,381
 mm) und etwa 55 mil (1,397 mm) dicke keramische Schicht ist, wobei die abreibbare Deckschicht ein Gemisch aus mit Cerdioxid teilstabilisiertem Zirkoniumdioxid und mit Cerdioxid vollständig stabilisiertem Zirkoniumdioxid aufweist.
 40

5. System nach Anspruch 1, 2 oder 3, bei dem die abreibbare Deckschicht (22) eine zwischen etwa 15 Mill (0,381
 mm) und etwa 55 mil (1,397 mm) dicke keramische Schicht ist, wobei die abreibbare Deckschicht ein Gemisch aus mit Magnesiumoxid teilstabilisiertem Zirkoniumdioxid und mit Magnesiumoxid vollständig stabilisiertem Zirkoniumdioxid aufweist.
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6. System nach Anspruch 1, 2 oder 3, bei dem die abreibbare Deckschicht (22) eine zwischen etwa 15 Mill (0,381
 mm) und etwa 55 mil (1,397 mm) dicke keramische Schicht ist, wobei die abreibbare Deckschicht ein Gemisch aus mit Yttriumoxid teilstabilisiertem Zirkoniumdioxid und mit Yttriumoxid vollständig stabilisiertem Zirkoniumdioxid aufweist.
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7. System nach Anspruch 1, 2 oder 3 bei dem die abreibbare Deckschicht (22) ein Material aufweist, das ausgewählt ist aus der Gruppe, die besteht aus mit Cerdioxid vollständig stabilisiertem Zirkoniumdioxid, mit Magnesiumoxid vollständig stabilisiertem Zirkoniumdioxid, mit Calciumoxid vollständig stabilisiertem Zirkoniumdioxid und mit Yttriumoxid vollständig stabilisiertem Zirkoniumdioxid.
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8. System nach Anspruch 7, bei dem die abreibbare Deckschicht (22) zu im wesentlichen 100 Gew.-% eine Zusammensetzung aus mit Yttriumoxid vollständig stabilisiertem Zirkoniumdioxid hat.

9. System nach Anspruch 1, 2 oder 3, bei dem die abreibbare Deckschicht (22) ein Gemisch aus mit 7 Gew.-% Yttriumoxid stabilisiertem Zirkoniumdioxid und mit 20 Gew.-% Yttriumoxid stabilisiertem Zirkoniumdioxid aufweist.
10. System nach Anspruch 1, bei dem
 - 5 die Basisbeschichtung-Grundlagenschicht zwischen etwa 5 mil (0,127 mm) und etwa 15 mil (0,381 mm) dick ist, wobei die Basisbeschichtung-Grundlagenschicht mit zwischen etwa 7 Gew.-% und etwa 12 Gew.-% Yttriumoxid stabilisiertes Zirkoniumdioxid mit einer Teilchengröße von etwa 50 Mikrometern (0,050 mm) im mittleren Durchmesser aufweist,
 - 10 die Varianz-Zwischenschicht zwischen etwa 3 mil (0,076 mm) und etwa 10 mil (0,254 mm) dick ist, wobei die Varianz-Zwischenschicht ein Zusammensetzungsgemisch aus der Basisbeschichtung-Grundlagenschicht und der abreibbaren Deckschicht aufweist, und
 - die abreibbare Deckschicht zwischen etwa 15 mil (0,381 mm) und etwa 55 mil (1,397 mm) dick ist, wobei die abreibbare Deckschicht mit 20 Gew.-% Yttriumoxid stabilisiertes Zirkoniumdioxid aufweist.
11. Strömungskanalsegment, aufweisend das segmentierte abreibbare keramische Beschichtungssystem gemäß einem der vorangehenden Ansprüche.
12. Dichtungssystem, aufweisend ein Strömungskanalsegment gemäß Anspruch 11 und ein in Wechselwirkung zusammenwirkendes Turbinenbauteil mit einer abreibenden Beschichtung auf einem Teil davon, so daß die abreibende Beschichtung zur Schaffung einer Abdichtung mit der segmentierten abreibbaren Beschichtung wechselwirken kann.
13. Verfahren zur Herstellung einer segmentierten abreibbaren keramischen Beschichtung, zu der eine Basisbeschichtung-Grundlagenschicht (18) eine Varianz-Zwischenschicht (20) und eine abreibbare Deckschicht (22) gehört, wobei das Verfahren aufweist:
 - 30 Aufbringen der Basisbeschichtung-Grundlagenschicht (18) auf ein mit MCrAlY bindungsbeschichtetes metallisches Substrat (12) unter Verwendung eines Sprühkopfes, der pulverförmiges Beschichtungsmaterial erhitzt und auf das metallische Substrat (12) treibt, wobei die Basisbeschichtung-Grundlagenschicht (18) eine Schicht aus einem Material aufweist, das ausgewählt ist aus der Gruppe, die besteht aus mit Cerdioxid stabilisiertem Zirkoniumdioxid, mit Magnesiumoxid stabilisiertem Zirkoniumdioxid, mit Calciumoxid stabilisiertem Zirkoniumdioxid, mit Yttriumoxid stabilisiertem Zirkoniumdioxid und Gemischen davon, wobei der Abstand zwischen dem Sprühkopf und der zu beschichtenden Oberfläche während der Herstellung der Basisbeschichtung-Grundlagenschicht verändert wird,
 - 35 Aufbringen der Varianz-Zwischenschicht (20) auf die Basisbeschichtung-Grundlagenschicht (18) unter Verwendung eines Sprühkopfes, der pulverförmiges Beschichtungsmaterial erhitzt und auf die Basisbeschichtung-Grundlagenschicht (18) treibt, wobei die Varianz-Zwischenschicht ein Zusammensetzungsgemisch aus der Basisbeschichtung-Grundlagenschicht und der abreibbaren Deckschicht aufweist, wobei der Abstand zwischen dem Kopf und der zu beschichtenden Oberfläche während der Herstellung der Varianz-Schicht konstant gehalten wird, und
 - 40 Aufbringen der abreibbaren Deckschicht (22) auf die Varianz-Zwischenschicht (20) unter Verwendung eines Sprühkopfes, der pulverförmiges Beschichtungsmaterial erhitzt und auf die Varianz-Zwischenschicht (20) treibt, wobei die abreibbare Deckschicht Zirkoniumdioxid aufweist, wobei der Abstand zwischen dem Kopf und der zu beschichtenden Oberfläche während der Herstellung der ; abreibbaren Deckschicht konstant gehalten wird, und wobei zwecks erhöhter Abreibbarkeit jede Schicht im wesentlichen aus Pulverteilchen, die vor der Abscheidung kugelförmig und hohl sind, besteht.
14. Verfahren nach Anspruch 13, bei dem die Basisbeschichtung-Grundlagenschicht in einer Dicke zwischen etwa 5 mil (0,127 mm) und etwa 15 mil (0,381 mm) und die Varianz-Schicht in einer Dicke zwischen etwa 3 mil (0,076 mm) und etwa 10 mil (0,254 mm) hergestellt wird.
15. Segmentiertes abreibbares keramisches Beschichtungssystem, bestehend im wesentlichen aus einer segmentierten abreibbaren keramischen Schicht und einem mit MCrAlY bindungsbeschichteten Substrat, wobei die abreibbare keramische Schicht ein Material aufweist, das ausgewählt ist aus der Gruppe, die besteht aus einem Material, das ein Zusammensetzungsgemisch aus mit Yttriumoxid teilstabilisiertem Zirkoniumoxid und mit Yttriumoxid vollständig stabilisiertem Zirkoniumoxid hat, einem Material, das ein Zusammensetzungsgemisch aus mit Cerdioxid teilstabilisiertem Zirkoniumdioxid und mit Cerdioxid vollständig stabilisiertem Zirkoniumdioxid hat, einem

Material, das ein Zusammensetzungsgemisch aus mit Magnesiumoxid teilstabilisiertem Zirkoniumdioxid und mit Magnesiumoxid vollständig stabilisiertem Zirkoniumdioxid hat, und einem Material, das ein Zusammensetzungsgemisch aus mit Calciumoxid teilstabilisiertem Zirkoniumdioxid und mit Calciumoxid vollständig stabilisiertem Zirkoniumdioxid hat, bei dem die abreibbare keramische Schicht auf dem mit MCrAlY bindungsbeschichteten metallischen Substrat angeordnet ist, wobei die Schicht zwecks erhöhter Abreibbarkeit im wesentlichen aus Pulverteilen, die vor der Abscheidung eine kugelförmige und hohle Morphologie haben, besteht.

16. Beschichtungssystem nach Anspruch 15, bei dem die keramische Schicht zwischen etwa 15 mil (0,381 mm) und etwa 55 mil (1,397 mm) dick ist.

Revendications

1. Système de revêtement céramique segmenté présentant une aptitude à l'abrasion, comprenant :

un subjectile métallique (12) ;
 une couche de fond (14) de MCrAlY sur le subjectile ;
 un revêtement céramique abradable segmenté (16) sur la couche de fond constituée de MCrAlY, ledit revêtement céramique abradable segmenté comprenant trois couches céramiques, qui comprennent une couche de fondation (18) formant couche de base, comprenant une couche d'un matériau choisi dans l'ensemble comprenant l'oxyde de zirconium stabilisé à l'oxyde de cérium, l'oxyde de zirconium stabilisé à l'oxyde de magnésium, l'oxyde de zirconium stabilisé à l'oxyde de calcium, l'oxyde de zirconium stabilisé à l'oxyde d'yttrium, et leurs mélanges ; une couche supérieure apte à l'abrasion (22) comprenant de l'oxyde de zirconium ; et une couche intermédiaire échelonnée (20), qui est un mélange en composition de la couche de fondation formant couche de base et de la couche de finition abradable, ladite couche intermédiaire échelonnée étant disposée entre la couche de formation formant couche de base et la couche supérieure apte à l'abrasion, ledit revêtement céramique abradable segmenté ayant une pluralité de microfissures verticales, dans lequel les trois couches céramiques sont constituées essentiellement de particules en poudre qui sont sphériques et creuses avant déposition, pour augmenter l'aptitude à l'abrasion.

2. Système selon la revendication 1, dans lequel la couche de fondation (18) formant couche de base est une couche céramique ayant une épaisseur d'environ 0,127 mm (5 mils) à environ 0,381 mm (15 mils), ladite couche de fondation formant couche de base étant choisie dans le groupe consistant en les matériaux contenant moins d'environ 12% en poids d'oxyde de zirconium stabilisé à l'oxyde de cérium, les matériaux contenant moins d'environ 12% en poids d'oxyde de zirconium stabilisé à l'oxyde de magnésium, les matériaux contenant moins d'environ 12% en poids d'oxyde de zirconium stabilisé à l'oxyde de calcium, les matériaux contenant moins d'environ 12% en poids d'oxyde de zirconium stabilisé à l'oxyde d'yttrium, et leurs mélanges.

3. Système selon la revendication 1, dans lequel la couche de fondation (18) formant couche de base comprend un mélange d'alumine et de moins d'environ 12% en poids d'oxyde de zirconium stabilisé à l'oxyde d'yttrium.

4. Système selon la revendication 1, 2 ou 3, dans lequel la couche de finition apte à l'abrasion (22) est une couche céramique ayant une épaisseur d'environ 0,381 mm (15 mils) à environ 1,397 mm (55 mils), ladite couche de finition apte à l'abrasion comprenant un mélange d'oxyde de zirconium partiellement stabilisé à l'oxyde de cérium et d'oxyde de zirconium entièrement stabilisé à l'oxyde de cérium.

5. Système selon la revendication 1, 2 ou 3, dans lequel la couche de finition apte à l'abrasion (22) est une couche céramique ayant une épaisseur d'environ 0,381 mm (15 mils) à environ 1,397 mm (55 mils), ladite couche de finition apte à l'abrasion comprenant un mélange d'oxyde de zirconium partiellement stabilisé à l'oxyde de magnésium et d'oxyde de zirconium entièrement stabilisé à l'oxyde de magnésium.

6. Système selon la revendication 1, 2 ou 3, dans lequel la couche de finition apte à l'abrasion (22) est une couche céramique ayant une épaisseur d'environ 0,381 mm (15 mils) à environ 1,397 mm (55 mils), ladite couche de finition apte à l'abrasion comprenant un mélange d'oxyde de zirconium partiellement stabilisé à l'oxyde d'yttrium et d'oxyde de zirconium entièrement stabilisé à l'oxyde d'yttrium.

7. Système selon la revendication 1, 2 ou 3, dans lequel la couche de finition apte à l'abrasion (22) comprend un matériau choisi dans le groupe constitué de l'oxyde de zirconium entièrement stabilisé à l'oxyde de cérium, de

l'oxyde de zirconium entièrement stabilisé à l'oxyde de magnésium, de l'oxyde de zirconium entièrement stabilisé à l'oxyde de calcium, et de l'oxyde de zirconium entièrement stabilisé à l'oxyde d'yttrium.

- 5 8. Système selon la revendication 7, dans lequel la couche de finition apte à l'abrasion (22) contient essentiellement 100% en poids d'oxyde de zirconium entièrement stabilisé à l'oxyde d'yttrium.
- 10 9. Système selon la revendication 1, 2 ou 3, dans lequel la couche de finition apte à l'abrasion (22) comprend un mélange de 7% en poids d'oxyde de zirconium stabilisé à l'oxyde d'yttrium et de 20% en poids d'oxyde de zirconium stabilisé à l'oxyde d'yttrium.
- 15 10. Système selon la revendication 1, dans lequel
la couche de fondation formant couche de base a une épaisseur d'environ 0,127 mm (5 mils) à environ 0,381 mm (15 mils), ladite couche de fondation formant couche de base contenant d'environ 7% en poids à environ 12% en poids d'oxyde de zirconium stabilisé à l'oxyde d'yttrium, avec une granulométrie, en diamètre moyen, d'environ 0,050 mm (50 µm) ;
la couche intermédiaire échelonnée a une épaisseur d'environ 0,076 mm (3 mils) à environ 0,254 mm (10 mils), ladite couche intermédiaire échelonnée comprenant un mélange de la couche de fondation formant couche de base et de la couche de finition abradable ; et
20 la couche de finition apte à l'abrasion a une épaisseur d'environ 0,381 mm (15 mils) à environ 1,397 mm (55 mils), ladite couche de finition apte à l'abrasion comprenant 20% en poids d'oxyde de zirconium stabilisé à l'oxyde d'yttrium.
- 25 11. Segment de gaine, comprenant le système de revêtement céramique apte à l'abrasion segmenté selon l'une quelconque des revendications précédentes.
- 30 12. Système d'étanchéité comprenant un segment de gaine selon la revendication 11 et un composant de turbine, en interaction par coopération, ayant un revêtement abrasif sur une portion de ce composant, tel que ledit revêtement abrasif puisse interagir avec ledit revêtement apte à l'abrasion segmenté pour assurer une étanchéité.
- 35 13. Procédé de production d'un revêtement céramique apte à l'abrasion segmenté comprenant une couche de fondation (18) formant couche de base, une couche intermédiaire échelonnée (20) et une couche de finition apte à l'abrasion (22), ledit procédé comprenant :
l'application de la couche de fondation (18) formant couche de base sur un support métallique (12) ayant
40 reçu une couche de fond constitué de MCrAlY, par utilisation d'un pistolet qui chauffe et projette un matériau de revêtement en poudre sur ledit support métallique (12), ladite couche de fondation (18) formant couche de base comprenant une couche d'un matériau choisi dans le groupe constitué par l'oxyde de zirconium stabilisé à l'oxyde de cérium, l'oxyde de zirconium stabilisé à l'oxyde de magnésium, l'oxyde de zirconium stabilisé à l'oxyde de calcium, l'oxyde de zirconium stabilisé à l'oxyde d'yttrium, et leurs mélanges, où la distance entre le pistolet et la surface à revêtir varie pendant la production de la couche de fondation formant couche de base, l'application de la couche intermédiaire échelonnée (20) sur la couche de fondation (18) formant couche de base par utilisation d'un pistolet qui chauffe et projette un matériau de revêtement en poudre sur ladite couche de fondation (18) formant couche de base, ladite couche intermédiaire échelonnée comprenant un mélange
45 de la couche de fondation formant couche de base et de la couche de finition abradable ; où la distance entre le pistolet et la surface à revêtir est maintenue constante pour la production de la couche échelonnée, et l'application de la couche de finition apte à l'abrasion (22) sur la couche intermédiaire échelonnée (20) par utilisation d'un pistolet qui chauffe et projette un matériau de revêtement en poudre sur ladite couche intermédiaire échelonnée (20), ladite couche de finition apte à l'abrasion comprenant de l'oxyde de zirconium, où
50 la distance entre le pistolet et la surface à revêtir est maintenue constante pendant la production de la couche de finition apte à l'abrasion, chaque couche étant constituée essentiellement de particules de poudre qui sont sphériques et creuses avant déposition, pour augmenter l'aptitude à l'abrasion.
- 55 14. Procédé selon la revendication (13) dans lequel la couche de fondation formant couche de base est produite sur une épaisseur d'environ 0,127 mm (5 mils) à environ 0,381 mm (15 mils), et la couche échelonnée est produite sur une épaisseur d'environ 0,076 mm (3 mils) à environ 0,254 mm (10 mils).
15. Système de revêtement céramique apte à l'abrasion segmenté constitué essentiellement d'une couche céramique

apte à l'abrasion segmentée et d'un subjectile ayant reçu une couche de fond constituée de MCrAlY, ladite couche
céramique apte à l'abrasion comprenant un matériau choisi dans le groupe consistant en un matériau composé
d'un mélange d'oxyde de zirconium partiellement stabilisé à l'oxyde d'yttrium et d'oxyde de zirconium entièrement
stabilisé à l'oxyde d'yttrium ; un matériau composé d'un mélange d'oxyde de zirconium partiellement stabilisé à
5 l'oxyde de cérium et d'oxyde de zirconium entièrement stabilisé à l'oxyde de cérium ; un matériau composé d'un
mélange d'oxyde de zirconium partiellement stabilisé à l'oxyde de magnésium et d'oxyde de zirconium entièrement
stabilisé à l'oxyde de magnésium ; et un matériau composé d'un mélange d'oxyde de zirconium partiellement
stabilisé à l'oxyde de calcium et d'oxyde de zirconium entièrement stabilisé à l'oxyde de calcium, où la couche
céramique apte à l'abrasion se trouve sur ledit subjectile métallique ayant reçu une couche de fond de MCrAlY, la
10 couche étant constituée essentiellement de particules de poudre ayant une morphologie sphérique ou creuse
avant déposition, pour augmenter l'aptitude à l'abrasion.

16. Système de revêtement selon la revendication 15, dans lequel la couche céramique a une épaisseur d'environ
0,381 mm (15 mils) à environ 1,397 mm (55 mils).

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fig. 1

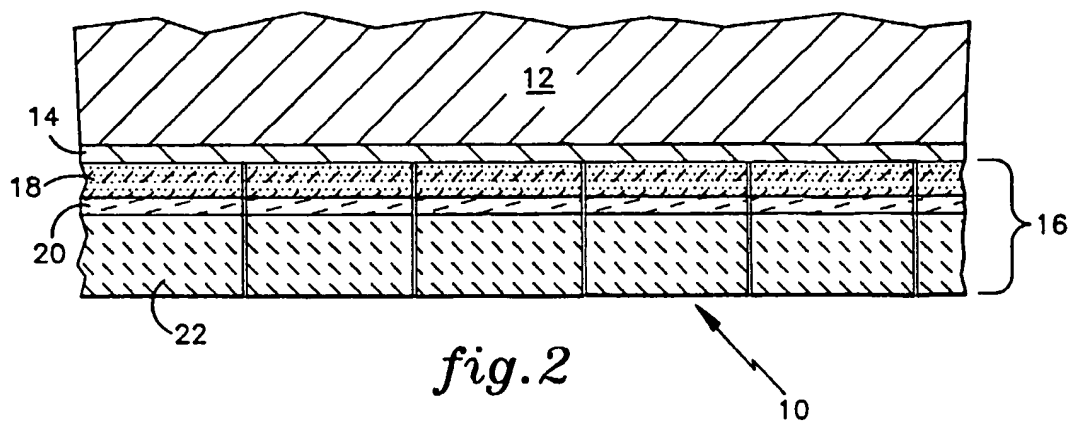
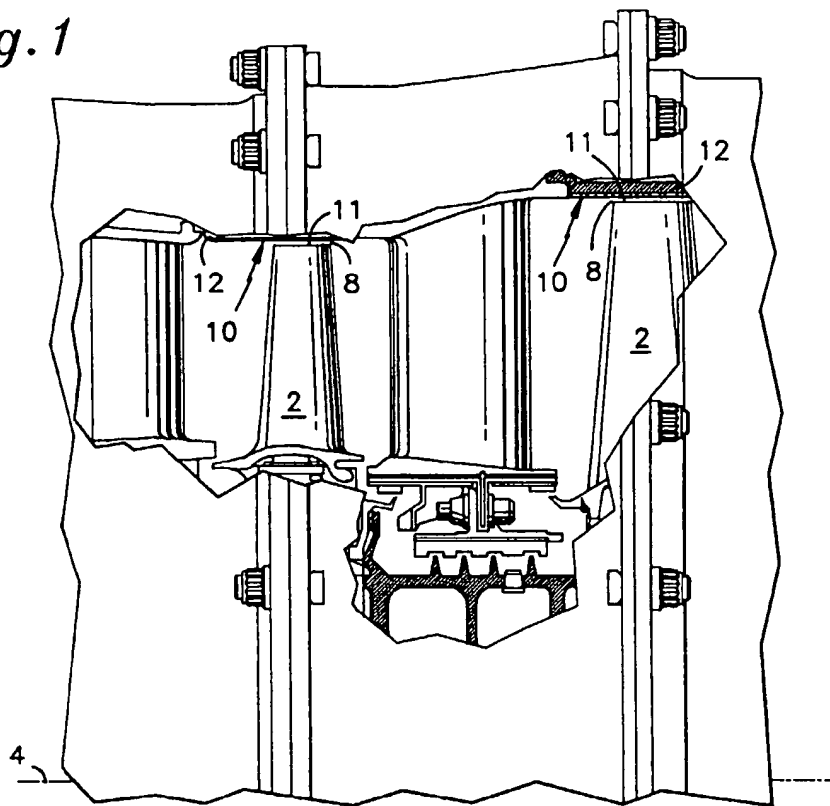


fig. 2